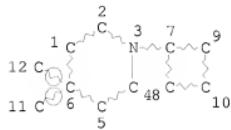


=> d 11
L1 HAS NO ANSWERS
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NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
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NUMBER OF NODES IS 12

STEREO ATTRIBUTES: NONE

=> s 11 ful
FULL SEARCH INITIATED 12:08:53 FILE 'REGISTRY'
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COST IN U.S. DOLLARS SINCE FILE TOTAL
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FILE COVERS 1907 - 17 Sep 2008 VOL 149 ISS 12
FILE LAST UPDATED: 16 Sep 2008 (20080916/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply.

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<http://www.cas.org/legal/infopolicy.html>

=> s 13
L4 12 L3

=> d bib 1-12

L4 ANSWER 1 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2008:353313 CAPLUS
DN 148:379624
TI Benzimidazolecarboxamide derivatives as bradykinin B1 receptor modulators and their preparation, pharmaceutical compositions and use in the treatment of inflammation and pain
IN Chandrasekhar, Jayaraman; Guo, Qin; Ihle, David C.; Ge, Ping; Wustrow, David J.; Chenard, Bertrand L.; Hodgetts, Kevin J.
PA Neurogen Corporation, USA
SO PCT Int. Appl., 82pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2008033739	A2	20080320	WO 2007-US77959	20070910
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW	RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CT, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
PRAI US 2006-825308P	P	20060912		
OS MARPAT 148:379624				

L4 ANSWER 2 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2007:1392027 CAPLUS
DN 148:54908
TI Preparation of spirocyclic sulfonamides and related compounds as modulators of bradykinin receptor activity
IN Hodgetts, Kevin J.; Ihle, David C.; Li, Guiying; Ge, Ping; Chenard, Bertrand L.; Wustrow, David J.
PA Neurogen Corporation, USA
SO PCT Int. Appl., 82pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2007140383	A2	20071206	WO 2007-US69918	20070530
WO 2007140383	A3	20080124		
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 PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN,
 TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
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 IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF,
 BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW,
 GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
 BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA

PRAI US 2006-803419P P 20060530
 OS MARPAT 148:54908

L4 ANSWER 3 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2007:1215844 CAPLUS

DN 147:486464

TI Preparation of 3-(piperazin-1-yl)cyclobut-3-ene-1,2-dione derivatives as antiviral agents

IN Bachand, Carol; Deon, Daniel H.; Ruediger, Edward H.

PA Bristol-Myers Squibb Company, USA

SO U.S. Pat. Appl. Publ., 38pp.

CODEN: USXXCO

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 20070249624	A1	20071025	US 2007-737354	20070419
	WO 2007127731	A1	20071108	WO 2007-US67302	20070424
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, IU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW				
RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				

PRAI US 2006-794699P P 20060425
 OS MARPAT 147:486464

L4 ANSWER 4 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2007:563421 CAPLUS

DN 147:9912

TI Preparation of diazacycloalkane-containing N,N-bis(1H-imidazol-2-ylmethyl)amines and related N,N-bis(heteroaryl methyl)amines as antagonists of chemokine receptor CXCR4

IN Kokubo, Masaya; Ochiai, Hiroshi; Takaoka, Yoshikazu; Shibayama, Shiro
PA Ono Pharmaceutical Co., Ltd., Japan

SO PCT Int. Appl., 332pp.

CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2007058322	A1	20070524	WO 2006-JP323015	20061117
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,				

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 MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO,
 RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT,
 TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
 RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
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 CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,
 GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
 KG, KZ, MD, RU, TJ, TM
 EP 1961744 A1 20080827 EP 2006-832893 20061117
 R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
 IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR
 PRAI JP 2005-334937 A 20051118
 JP 2006-49378 A 20060224
 WO 2006-JE323015 W 20061117
 OS MARPAT 147:9912
 RE.CNT 42 THERE ARE 42 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 5 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
 AN 2007:203199 CAPLUS
 DN 146:274233
 TI Preparation of spiro-fused piperidines as modulators of muscarinic receptors
 IN Makings, Lewis R.; Garcia-Guzman Blanco, Miguel; Hurley, Dennis J.; Drutu, Ioana; Raffai, Gabriel; Bergeron, Daniele M.; Nakatani, Akiko; Termin, Andreas P.; Silina, Alina
 PA USA
 SO U.S. Pat. Appl. Publ., 500pp., Cont.-in-part of U.S. Ser. No. 208,386.
 CODEN: USXXCO
 DT Patent
 LA English
 FAN.CNT 2

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 20070043023	A1	20070222	US 2006-359960	20060222
US 20060079505	A1	20060413	US 2005-208386	20050819
PRAI US 2004-602731P	P	20040819		
US 2005-208386	A2	20050819		
OS MARPAT 146:274233				

L4 ANSWER 6 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
 AN 2006:693858 CAPLUS
 DN 145:201982
 TI 4-Substituted-8-(1-phenyl-cyclohexyl)-2,8-diaza-spiro[4.5]decan-1-one as a novel class of highly selective GlyT1 inhibitors with superior pharmacological and pharmacokinetic parameters
 AU Alberati, Daniela; Hainzl, Dominik; Jolidon, Synesse; Kurt, Anke; Pinard, Emmanuel; Thomas, Andrew W.; Zimmerli, Daniel
 CS Discovery Biology, Pharmaceuticals Division, F. Hoffmann-La Roche Ltd., Basel, CH-4070, Switz.
 SO Bioorganic & Medicinal Chemistry Letters (2006), 16(16), 4321-4325
 CODEN: BMCLE8; ISSN: 0960-894X
 PB Elsevier B.V.
 DT Journal
 LA English
 OS CASREACT 145:201982
 RE.CNT 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 7 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2006:193777 CAPLUS
DN 144:274274
TI Preparation of basic group-containing heterocyclic compounds as CXCR4 antagonists

IN Kokubo, Masaya; Takaoka, Yoshikazu; Shibayama, Shiro
PA Ono Pharmaceutical Co., Ltd, Japan
SO PCT Int. Appl., 305 pp.
CODEN: PIXXD2

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006022454	A1	20060302	WO 2005-JP16066	20050826
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	EP 1790639	A1	20070530	EP 2005-776646	20050826
	R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				
	US 20080009495	A1	20080110	US 2007-661270	20070227
PRAI	JP 2004-248431	A	20040827		
	JP 2005-100039	A	20050330		
	JP 2005-190741	A	20050629		
	WO 2005-JP16066	W	20050826		

OS MARPAT 144:274274

RE.CNT 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 8 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
AN 2006:15778 CAPLUS
DN 144:108321

TI Preparation of amino cyclopentyl heterocyclic and carbocyclic modulators
of chemokine receptor activity

IN Yang, Lihu; Lin, Songnian; Morriello, Gregori; Guo, Liangqin; Zhou,
Changyou

PA Merck & Co., Inc., USA
SO PCT Int. Appl., 141 pp.
CODEN: PIXXD2

DT Patent
LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006001958	A2	20060105	WO 2005-US17836	20050520
	WO 2006001958	A3	20060824		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK,				

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 ZA, ZM, ZW
 RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
 IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF,
 CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM,
 KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG,
 KZ, MD, RU, TJ, TM
 AU 2005257859 A1 20060105 AU 2005-257859 20050520
 CA 2567851 A1 20060105 CA 2005-2567851 20050520
 EP 1753740 A2 20070221 EP 2005-785401 20050520
 R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
 IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA,
 HR, LV, MK, YU
 CN 1956975 A 20070502 CN 2005-80016231 20050520
 JP 2008502719 T 20080131 JP 2007-527495 20050520
 IN 2006DN06362 A 20070831 IN 2006-DN6362 20061030
 PRAI US 2004-573625P P 20040521
 WO 2005-US17836 W 20050520
 OS MARPAT 144:108321

L4 ANSWER 9 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
 AN 2005:612285 CAPLUS
 DN 143:133293
 TI Preparation of spiroindoline and spiroisoquinoline compounds as Mas receptor ligands
 IN Boatman, Douglas P.; Adams, John W.; Moody, Jeanne V.; Babych, Eric D.; Schrader, Thomas O.
 PA Arena Pharmaceuticals, Inc., USA
 SO PCT Int. Appl., 224 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2005063745	A2	20050714	WO 2004-US43609	20041222
WO 2005063745	A3	20060316		
WO 2005063745	A9	20070201		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, SM RW: BN, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2004309419	A1	20050714	AU 2004-309419	20041222
CA 2546147	A1	20050714	CA 2004-2546147	20041222
EP 1716148	A2	20061102	EP 2004-815636	20041222
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, BA, HR, IS, YU				
CN 1972944	A	20070530	CN 2004-80038166	20041222
JP 2007516298	T	20070621	JP 2006-547461	20041222
IN 2006KN02015	A	20070518	IN 2006-KN2015	20060718
US 20070254903	A1	20071101	US 2007-583839	20070308
PRAI US 2003-532546P	P	20031223		
US 2004-539554P	P	20040126		

US 2004-565251P P 20040423
 WO 2004-US43609 W 20041222
 OS MARPAT 143:133293

L4 ANSWER 10 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
 AN 2004:1124588 CAPLUS
 DN 142:69197
 TI CCR-2 antagonists for treatment of neuropathic pain
 IN Abbadie, Catherine; Lindia, Jill Ann; Wang, Hao
 PA Merck & Co., Inc., USA
 SO PCT Int. Appl., 304 pp.
 CODEN: PIXXD2

DT Patent
 LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004110376	A2	20041223	WO 2004-US17499	20040602
	WO 2004110376	A3	20050224		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, US, UZ, VC, VN, YU, ZA, ZM, ZW, RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	US 20060205761	A1	20060914	US 2005-559701	20051206
PRAI	US 2003-476391P	P	2003030606		
	US 2003-531637P	P	20031222		
	WO 2004-US17499	W	20040602		
OS	MARPAT 142:69197				

L4 ANSWER 11 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
 AN 2004:802715 CAPLUS
 DN 141:314157
 TI Preparation of amino cyclobutylamide modulators of chemokine receptor activity

IN Jiao, Richard; Yang, Lihu
 PA Merck & Co. Inc., USA
 SO PCT Int. Appl., 108 pp.
 CODEN: PIXXD2

DT Patent
 LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004082682	A1	20040930	WO 2004-US7792	20040315
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN,				

TD, TG
 AU 2004222336 A1 20040930 AU 2004-222336 20040315
 CA 2519220 A1 20040930 CA 2004-2519220 20040315
 EP 1617841 A1 20060125 EP 2004-720791 20040315
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
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 CN 1787818 A 20060614 CN 2004-80013143 20040315
 JP 2006520783 T 20060914 JP 2006-507176 20040315
 IN 2005DN03929 A 20070824 IN 2005-DN3929 20050902
 US 20060211722 A1 20060921 US 2005-549739 20050919
 PRAI US 2003-456047P P 20030318
 WO 2004-US7792 A 20040315
 OS MARPAT 141:314157
 RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 12 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
 AN 2001:247141 CAPLUS
 DN 134:266327
 TI Preparation of spiro[bicyclic-azacycloalkyl and -cycloalkyl] derivatives
 and uses as alpha 1a adrenergic receptor antagonists
 IN Hoffman, Jacob M.
 PA Merck & Co., Inc., USA
 SO PCT Int. Appl., 140 PP.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

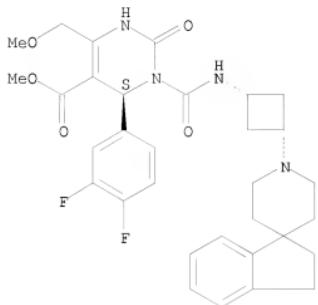
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2001022919	A2	20010405	WO 2000-US26387	20000926
WO 2001022919	A3	20020711		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
AU 2000077166	A	20010430	AU 2000-77166	20000926
PRAI US 1999-157207P	P	19990930		
WO 2000-US26387	W	20000926		
OS MARPAT 134:266327				

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L4 ANSWER 12 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN
 IT 332187-08-3P 332187-09-4P 332187-37-8P
 332187-38-9P
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological
 study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use);
 BIOL (Biological study); PREP (Preparation); USES (Uses)
 (preparation of spirobicyclazacycloalkyl and cycloalkyl derivs. and uses
 as alpha a adrenergic receptor antagonists)
 RN 332187-08-3 CAPLUS
 CN 5-Pyrimidinecarboxylic acid, 6-(3,4-difluorophenyl)-1-[[[cis-3-(2,3-
 dihydrospiro[1H-indene-1,4'-piperidin]-1'-yl)cyclobutyl]amino]carbonyl]-
 1,2,3,6-tetrahydro-4-(methoxymethyl)-2-oxo-, methyl ester, hydrochloride

(1:1), (6S)- (CA INDEX NAME)

Absolute stereochemistry.

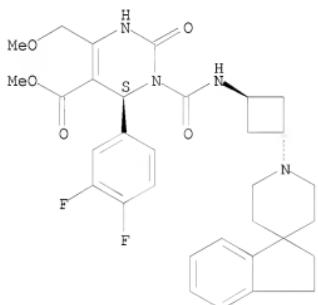


● HCl

RN 332187-09-4 CAPLUS

CN 5-Pyrimidinecarboxylic acid, 6-(3,4-difluorophenyl)-1-[(trans-3-(2,3-dihydrospiro[1H-indene-1,4'-piperidin]-1'-yl)cyclobutyl)amino]carbonyl]-1,2,3,6-tetrahydro-4-(methoxymethyl)-2-oxo-, methyl ester, hydrochloride
(1:1), (6S)- (CA INDEX NAME)

Absolute stereochemistry.

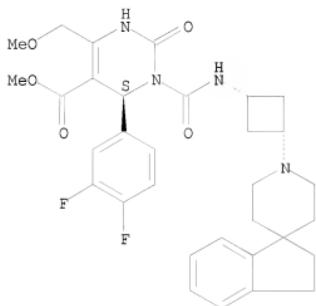


● HCl

RN 332187-37-8 CAPLUS

CN 5-Pyrimidinecarboxylic acid, 6-(3,4-difluorophenyl)-1-[[[cis-3-(2,3-dihydrospiro[1H-indene-1,4'-piperidin]-1'-yl)cyclobutyl]amino]carbonyl]-1,2,3,6-tetrahydro-4-(methoxymethyl)-2-oxo-, methyl ester, (6S)- (CA INDEX NAME)

Absolute stereochemistry.



RN 332187-38-9 CAPLUS

CN 5-Pyrimidinecarboxylic acid, 6-(3,4-difluorophenyl)-1-[[[trans-3-(2,3-dihydrospiro[1H-indene-1,4'-piperidin]-1'-yl)cyclobutyl]amino]carbonyl]-1,2,3,6-tetrahydro-4-(methoxymethyl)-2-oxo-, methyl ester, (6S)- (CA INDEX NAME)

Absolute stereochemistry.

